

What is Claimed is

1. A method for treating the effects of low LDL-cholesterol values in the brain on cognitive performance or other central nervous system functions by modulating nicotinic receptors by administering an effective amount of a nicotinic allosteric potentiator, an acetylcholinesterase inhibitor, nicotine, a nicotinic agonist or a mixture thereof to a patient in need of such modulation.
2. A method as claimed in claim 1 wherein said low cholesterol values are values of less than 109 mg/dl LDL-cholesterol.
3. A method as claimed in claim 1 wherein said low cholesterol values are the result of treatment with HMG-CoA reductase inhibitors.
4. A method as claimed in claim 1 wherein said modulation of nicotinic receptors is effected by administering an effective amount of a galanthamine or lycoramine analog to a patient in need of such modulation.
5. A method as claimed in claim 2 wherein said modulation of nicotinic receptors is effected by administering an effective amount of a galanthamine or lycoramine analog to a patient in need of such modulation.

6. A method as claimed in claim 3 wherein said modulation of nicotinic receptors is effected by administering an effective amount of a galanthamine or lycoramine analog to a patient in need of such modulation.

7. A method as claimed in claim 4 wherein said analog of galanthamine or lycoramine is one wherein at least one of the methoxy, hydroxy or methyl groups of galanthamine or lycoramine is replaced as follows:

the methoxy group by another alkoxy group of from one to six carbon atoms, a hydroxy group, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group or a trialkylsilyloxy group;

the hydroxy group by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group;

the N-methyl group by hydrogen, alkyl, benzyl, cyclopropylmethyl group or a substituted or unsubstituted benzoyloxy group.

8. A method as claimed in claim 5 wherein said analog of galanthamine or lycoramine is one wherein at least one of the methoxy, hydroxy or methyl groups of galanthamine or lycoramine is replaced as follows:

the methoxy group by another alkoxy group of from one to six carbon atoms, a hydroxy group, hydrogen, an alkanoyloxy group, a benzoyloxy or

substituted benzoyloxy group, a carbonate group or a carbamate group or a trialkylsilyloxy group;

the hydroxy group by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group;

the N-methyl group by hydrogen, alkyl, benzyl, cyclopropylmethyl group or a substituted or unsubstituted benzoyloxy group.

9. A method as claimed in claim 6 wherein said analog of galanthamine or lycoramine is one wherein at least one of the methoxy, hydroxy or methyl groups of galanthamine or lycoramine is replaced as follows:

the methoxy group by another alkoxy group of from one to six carbon atoms, a hydroxy group, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group or a trialkylsilyloxy group;

the hydroxy group by an alkoxy group of from one to six carbon atoms, hydrogen, an alkanoyloxy group, a benzoyloxy or substituted benzoyloxy group, a carbonate group or a carbamate group;

the N-methyl group by hydrogen, alkyl, benzyl, cyclopropylmethyl group or a substituted or unsubstituted benzoyloxy group.

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10. A method as claimed in claim 4 wherein any alkanoyloxy, carbamate and carbonate group present contains up to ten carbon atoms.
11. A method as claimed in claim 5 wherein any alkanoyloxy, carbamate and carbonate group present contains up to ten carbon atoms.
12. A method as claimed in claim 6 wherein any alkanoyloxy, carbamate and carbonate group present contains up to ten carbon atoms.
13. A method as claimed in claim 4 wherein any of said alkanoyloxy, carbamate or carbonate group comprises an alkyl or alkoxy group of from 1 to 6 carbon atoms optionally substituted by one or more halo groups.
14. A method as claimed in claim 5 wherein any of said alkanoyloxy, carbamate or carbonate group comprises an alkyl or alkoxy group of from 1 to 6 carbon atoms optionally substituted by one or more halo groups.
15. A method as claimed in claim 6 wherein any of said alkanoyloxy, carbamate or carbonate group comprises an alkyl or alkoxy group of from 1 to 6 carbon atoms optionally substituted by one or more halo groups.
16. A method as claimed in claim 4 wherein said analog is a galanthamine analog.

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17. A method as claimed in claim 5 wherein said analog is a galanthamine analog.
18. A method as claimed in claim 6 wherein said analog is a galanthamine analog.
19. A method as claimed in claim 4 wherein said analog is a lycoramine analog.
20. A method as claimed in claim 5 wherein said analog is a lycoramine analog.
21. A method as claimed in claim 6 wherein said analog is a lycoramine analog.
22. A method as claimed in claim 4 wherein said analog is an n-butyl carbamate.
23. A method as claimed in claim 5 wherein said analog is an n-butyl carbamate.
24. A method as claimed in claim 6 wherein said analog is an n-butyl carbamate.
25. A method as claimed in claim 4 wherein the methoxy group of galanthamine or lycoramine is replaced by a hydrogen, hydroxy or alkoxy group of from two to six carbon atoms or an acyloxy group or a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.
26. A method as claimed in claim 5 wherein the methoxy group of galanthamine or lycoramine is replaced by a hydrogen, hydroxy or alkoxy group of from two to

six carbon atoms or an acyloxy group or a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.

27. A method as claimed in claim 6 wherein the methoxy group of galanthamine or lycoramine is replaced by a hydrogen, hydroxy or alkoxy group of from two to six carbon atoms or an acyloxy group or a mono or dialkyl carbamate or carbonate group wherein the alkyl groups contain from 1 to 8 carbon atoms.
28. A method as claimed in claim 4 wherein the hydroxy group of galanthamine or lycoramine is replaced by an alkoxy group of from one to six carbon atoms, hydrogen, an acyloxy group, a carbonate group or a carbamate group which may be a mono or dialkyl or an aryl carbamate or carbonate wherein the alkyl groups contain from 1 to 8 carbon atoms.
29. A method as claimed in claim 5 wherein the hydroxy group of galanthamine or lycoramine is replaced by an alkoxy group of from one to six carbon atoms, hydrogen, an acyloxy group, a carbonate group or a carbamate group which may be a mono or dialkyl or an aryl carbamate or carbonate wherein the alkyl groups contain from 1 to 8 carbon atoms.
30. A method as claimed in claim 6 wherein the hydroxy group of galanthamine or lycoramine is replaced by an alkoxy group of from one to six carbon atoms, hydrogen, an acyloxy group, a carbonate group or a carbamate group which may

group of two to six carbon atoms or a carbonate of from one to six carbon atoms an alkyl carbonate of one to six carbon atoms in its alkyl group or an aryl carbonate.

35. A method as claimed in claim 5 wherein the compound employed is one wherein the methoxy group of galanthamine or lycoramine is replaced by an alkoxy group of two to six carbon atoms or a carbonate of from one to six carbon atoms an alkyl carbonate of one to six carbon atoms in its alkyl group or an aryl carbonate.
36. A method as claimed in claim 6 wherein the compound employed is one wherein the methoxy group of galanthamine or lycoramine is replaced by an alkoxy group of two to six carbon atoms or a carbonate of from one to six carbon atoms an alkyl carbonate of one to six carbon atoms in its alkyl group or an aryl carbonate.
37. A method for treating neuromuscular dysfunction resulting from use of HMG-CoA reductase inhibitors by modulating nicotinic receptors by administering an effective amount of a nicotinic allosteric potentiator, nicotine, a nicotinic agonist or a mixture thereof to a patient in need of such modulation.